

AUTHORS: Levadnyy, G.Ye., Brodetskiy, Ye.S. 6-58-5-14/17

TITLE: The Calculation of Standards for Turnover Materials Within the Plan of an Air-Geodetical Undertaking (Raschet normativa oborotnykh sredstv v plane aerogeodezicheskogo predpriyatiya)

PERIODICAL: Geodeziya i Kartografiya, 1958, Nr 5, pp. 66-75 (USSR)

ABSTRACT: Working materials are subdivided into basic- and turnover materials. The basic principles in planning the requirements of an enterprise or firm with respect to turnover materials were laid down by the decree issued by the work- and defense council dated July 23, 1931, which is still in force. In the present case the experience gathered by an air-geodetical enterprise is generalized, and on the basis of data obtained for a number of years, a model-calculation of standards for turnover materials was set up. It is shown that, within the structure of assets to be standardized, stock of low value and objects subjected to a high degree of wear occupy the most important place (more than half of all assets). Basic- and auxiliary material range second. The second characteristic feature of the existing structure is the fact that all assets to be standardized are investments made in the turnover-production fund.

Card 1/2

The Calculation of Standards for Turnover Materials
Within the Plan of an Air-Geodetical Undertaking

6-58-5-14/17

Calculation is not based upon the extent to which plans are realized, but upon the volume of production, the structure of production costs as well as upon other indices connected herewith. The problem of the required level for individual types of stocks according to their importance for production and according to the specific weight within the structure of assets to be standardized are dealt with in detail. On the basis of these analytical data a workable method of calculating standards for the materials turned over within the plan of an air-geodetical enterprise is employed. There are 8 tables

1. Geophysical surveying---Materials
2. Materials---Standards
3. Mathematics

Card 2/2

3(4)

SOV/6-59-4-13/20

AUTHORS: Levadnyy, G. Ye., Brodetskiy, Ye. S.

TITLE: Organization of the Economic Book-keeping in an Aerogeodetic Enterprise (Organizatsiya khozyaystvennogo rascheta v aerogeodezicheskoy predpriyatii)

PERIODICAL: Geodeziya i kartografiya, 1959, Nr 4, pp 40-49 (USSR)

ABSTRACT: The editors invite the heads of aerogeodetic enterprises, heads of field squads, and other readers to take part in the discussion of the problems put forward here. An attempt is made here to generalize the practice of economic book-keeping. The first chapter investigates the basic conditions for economic book-keeping in aerogeodetic services. The introduction of regular book-keeping within the individual departments of the enterprise : the squads, shops, brigades and groups, is considered most important. The form of book-keeping in the individual departments within an enterprise greatly differs from the book-keeping of the enterprise itself. The basis for each such subsection of the enterprise is the corresponding plan. These plans should provide for book-keeping indices. Their characteristics should be considered in introducing the book-keeping in the auxiliary services. It is point-

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Organization of the Economic Book-keeping in an Aerogeodetic Enterprise

ed out in detail what should be considered in determining the indices for transportation by car, for transportation by horse carriages and for the supply of timber. The second chapter investigates the relationship between the individual subsections of the enterprise with respect to economic book-keeping. The third chapter explains the principle of material interest. The leading personnel should be paid according to the time-premium-wage system. The premium should depend, however, on the amount of saving. The premiums should be paid from the means of the premium fund of the corresponding department, not from the means of the fund of the enterprise. The fourth chapter deals with evidence and accounting. It is pointed out that up to date no instructions for the planning, evidence, and calculation of cost of production are available for the aerogeodetic services. The fifth chapter concerns financing. Two examples are given: a calculation of financing for a squad for one year, and a calculation for one month. There are 3 tables.

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3(4)

AUTHORS: Sirenko, I. F., Brodetakiy, Ye. S. SOV/6-59-8-17/27

TITLE: Competition for the Title of Brigades of Communist Work at the Ukrainian Aerogeodetic Enterprise (Sorevnovaniye za zvaniye brigad kommunisticheskogo truda v Ukrainskom aerogeodeticheskom predpriyatii)

PERIODICAL: Geodeziya i kartografiya, 1959, Nr 8, pp 64-65 (USSR)

ABSTRACT: Several months have passed since the competition first started at the above enterprise. A public meeting of Komsomol members was held on June 4. The young people of the enterprise attended the meeting in order to hear what the situation of the competition was. The following persons took the floor: Ira Shishova, Secretary of the Komsomol Organization, opened the meeting. The next speaker was Engineer Nelli Rytsk (other members of the brigade are Sinitskaya, Nikolayeva, Spareva, Protasevich), followed by Galya Zozulya, in whose brigade there is also Ludmila Kucheruk (the brigade cooperates closely with the brigade of Mayya Karpyuk), and Tanya Kissa, in whose brigade are Nadya Polozova, Tamara Tikhonova, Ira Logvinenko,

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Competition for the Title of Brigades of Communist
Work at the Ukraina Aerogeodetic Enterprise

SOV/6-59-8-17/27

Lyuda Stepanenko, Raya Kotel'nikova. Another speaker was
Nina Boykova, a worker from the candy factory imeni Karla
Marksa (imeni Karl Marx), who discussed the experiences
gathered in her brigade. - The meeting agreed to fulfil the
Seven-Year-Plan in six years.

Card 2/2

BRODFELD, B.

A method of static computation of elastic dam soles rigidly fixed on the contour. p. 79.

HIDROTEHNICA. (Asociatia Stiintifica a Inginerilor si Tehnicienilor din Romina) Bucuresti, Rumania, Vol. 4, no. 3, Mar. 1959.

Monthly list of East European Accessions (EEAI) LC Vol. 8, No. 9, ^{Sept.} 1959.

Uncl.

BRODFELD, B.

TECHNOLOGY

Periodicals: HIDROTEHNICA. Vol. 3, no. 10, Oct. 1958

BRODFELD, B. Some aspects of computing the foundation rafts of decantation apparatus. p. 375

Monthly List of East European Accessions (EEAI) LC, Vol. 8, No. 2,
February 1959, Unclass.

BRODI, S., nauchnyy sotrudnik; KOROLYUK, V., kand. fiz.- matem. nauk

Mass servicing as a mathematical problem . Nauka i zhyttia 13
no.10:6-7 N '63. (MIRA 16:12)

1. Institut matematiki AN UkrSSR (for Brodi).

16 (1)

SOV/21-59-6-2/27

AUTHOR: Brodi, S. M.

TITLE: On an Integral-Differential Equation for Systems
with τ -Waiting

PERIODICAL: Dopovidi Akademii Nauk Ukrain's'koi RSR, 1959, Nr 6,
pp 571 - 573 (USSR)

ABSTRACT: The author examines the simplest flow of demands on the service of one device. The time of service is a random value of exponential distribution. The demand entering the system is served immediately if the device is not engaged, and has to wait in line if the device is engaged. The demand waits for service no longer than the time τ , after which it is lost. Integro-differential equations for the time of waiting for service are formulated by the author. The system with waiting is a system where the demand waits only a limited time, and can be served only when the device is free, or will be free within a time less than $\tau = \text{const}$ ($\tau > 0$). Designations used in the calculations are standard mathematical.

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SOV/21-59-6-2/27
On an Integral-Differential Equation for Systems with τ -Waiting

There are 3 references, 1 of which is American,
1 Hungarian and 1 Soviet

ASSOCIATION: Institut matematiki AN UkrSSR (Institute of Mathematics of
the AS UkrSSR)

PRESENTED: By B. V. Gnedenko, Member, AS UkrSSR

SUBMITTED: December 25, 1958

Card 2/2

BRODI, S. M.

PLATE I BOX EMBLEM 90/1981

Sovetskimiye po teorii veroyatnostey i matematicheskoy statistike, Yerevan, 1958
 Tredy Mesnyatsnogo sovetskikh po teorii veroyatnostey i matematicheskoy
 statistike, Yerevan, 1958. 12-24. Yerevan: Izdatel'stvo Konferentsii po
 Teorii Veroyatnosti i Matematicheskoy Statistike. Held in Yerevan 19-25
 September, 1958. Translations: Yerevan, Izdatel'stvo AN SSR, 1960. 291 p.
 Entirely printed. 2,500 copies printed.

Sponsoring Agency: Akademiya nauk Armianskoy SSR.

Editorial Staff: G.A. Amartsyan, B.Y. Gekht, Yu.B. Dymkin, Yu.V. Klimik and
 S. Kh. Tsamiryan; Ed. of Publishing House: A.G. Shimi; Tech. Ed.: M.A. Kopylov.

NOTE: The book is intended for mathematicians.

CONTENTS: The book contains 11 articles submitted to the Conference and dealing with
 the theory of probability and mathematical statistics. Some of the articles are
 the papers read at the Conference and edited for publication, while others outline
 the theses of papers which appeared or are scheduled to appear, wholly or in
 part, in other publications; in some cases, such publications are quoted. A
 list of the papers whose contents were published elsewhere is included and the
 places of publication are indicated. Individual articles examine theories of
 mass service, spectral instruments, numbers, games, and certain functions, and
 discuss the theorems of Shannon, Markov's chains, and certain processes, quan-
 titles, and functions. Such items as the method of least squares, the stochastic
 Markov's and diffusion processes, measures and their applications, a scheme of
 Bernoulli experiments, Markov-type random fields, visible distribution of stars,
 Brownian motion, capacity of radio channels, and defective products are con-
 sidered. No personalities are mentioned. References accompany some of the
 articles.

Belman, R.P. Asymptotic Cardinality of Some Separable Criteria Concerning Displacement. (Theses)	98
Bernshteyn, O.Y. On Matrices Coefficient of Correlation. (Theses)	101
Bogdanov, A.A. New Results Concerning Independent Statistics. (Theses)	103
Bolshakov, O.K. On the Theory of the Method of Least Squares When Weights are Unknown	106
Amartsyan, G.A. On Quantity of Information About an Unknown Probability In the Scheme of Bernoulli's Experiments	112
Tsamiryan, S.K. On the Statistical Criterion, χ^2 , as Applied to the Problem of Two Samples	121
Amartsyan, Y.A. On Fluctuations in the Visible Distribution of Stars	129
Brod, S.M. On One Problem in the Theory of Mass Service Evidently, I.I. On the Restoration of Additive Type of Distribution by The Sequence of Series of Independent Observations	133
Klov, J.M. Random Quantities of Noncompact Semigroups. (Theses)	140
Kobayashi, I.P. Ts.Y. Klimik, and B.Y. Ushakov. Some New Results in the Probabilistic Theory of Numbers, and Simulation of Brownian Motion. (Theses)	162
Pokrovskiy, E.L., I.I. Khurgin, and B.S. Tyupakov. Approximate Compu- tation of the Carrying Capacity of Radio Channels with Random Parameters	164
Korotkiy, Kh.B. Distribution of the Number, X , of Defective Products in Lots	172
Shafin, L.A. On Theoretical Informational Approach to the Theory of Spectral Instruments	177
Romanovskiy, I.B. On Probability Problems Leading to Dynamic Programming	205

Card 6/8

16.6100

30851
S/044/61/000/008/028/039
C111/C333

AUTHOR: Brodi, S. M.

TITLE: On a problem of the theory of queuing

PERIODICAL: Referativnyy zhurnal, Matematika, no. 8, 1961, 26
abstract 8V179. ("Tr. Vses. soveshchaniya po teorii
veroyatnostey i matem. statistiki", 1958, Yerevan, AN
Arm SSR, 1960, 143-147)

TEXT: Assume that a stationary Poisson input flow with the
parameter $\alpha > 0$ is given. The service time is distributed according to
an exponential law with the parameter $\beta > 0$. The waiting time
 $\xi^{(\tau)}(t)$ of a visitor arriving at the moment t is bounded by the constant
 $\tau > 0$. Let

$$F^{(\tau)}(t, x) \equiv P \{ \xi^{(\tau)}(t) \leq x \mid \xi^{(\tau)}(t) < \tau \}$$

Then $F^{(\tau)}(t, x)$ satisfies the equation

$$\frac{\partial F^{(\tau)}(t, x)}{\partial t} = \frac{\partial F^{(\tau)}(t, x)}{\partial x} -$$

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C111/C333

On a problem of the theory of . . .

$$-\alpha \left[F^{(\tau)}(t, x) - \int_0^x (1 - e^{-\beta(x-y)}) dF^{(\tau)}(t, y) \right]$$

If $\alpha/\beta < 1$, then we have

$$\frac{dF_{\tau}^{*}(x)}{dx} = \alpha \left\{ F_{\tau}^{*}(x) - \int_0^x [1 - e^{-\beta(x-y)}] dF_{\tau}^{*}(y) \right\}$$

for $F_{\tau}^{*}(x) \equiv \lim_{t \rightarrow \infty} F^{(\tau)}(t, x)$. Let

$$\tilde{F}_{\tau}(p) = \int_0^{\infty} e^{-px} dF_{\tau}^{*}(x) \quad (\operatorname{Re}(p) > 0).$$

Then it holds

Card 2/3

On a problem of the theory of . . .

³⁰⁸⁵¹
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$$\tilde{F}_{\tau}(P) = \left[F_{\tau}(0) - e^{-\beta\tau} (1 - F(\tau)) \right] / \left[1 - \frac{\alpha(P+\beta)e^{-\tau(P+\beta)}}{P(P+\beta)} \right].$$

Note of the reviewer: The problem is considered without the restriction $\xi^{(\tau)}(t) < \tau$ in another paper of the author (Dokl. AN Ukr SSR, 1959, 6, 571-573).

[Abstracter's note: Complete translation.]

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Card 3/3

44206

16.6100

S/021/62/000/011/002/013
D251/D308

AUTHOR: Brodi, S. M.

TITLE: Unilinear system with τ -waiting in the case of an Erlang incoming stream

PERIODICAL: Akademiya nauk Ukrayins'koyi RSR. Dopovidi, no. 11, 1962, 1425-1428

TEXT: A system of service in which a client must wait for a finite time $\tau > 0$ is called a system with τ -waiting. The author considers the problem where clients arrive at times $\{t_i\}$ ($i = 1, 2, \dots$) such that the values of $t_i - t_{i-1}$ form an Erlang distribution and the service times $\{y_i\}$ ($i = 1, 2, \dots$) of the respective clients are independent and equally distributed random quantities. The simplified case $G_k(x) = [G_0(x)]^k$ is considered, where $P\{t_i - t_{i-1} \leq x\} = G_k(x)$ and by considering the random process $\eta_\tau(t)$ -

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Unilinear system with ...

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the waiting time of a client joining the system at a time t , and writing $P\{\eta(t) \leq x\} = F(t, x) = \sum_{j=1}^k F_j(t, x)$, a system of differential equations is obtained which must be satisfied by $F_j^*(x) = \lim_{t \rightarrow \infty} F_j(t, x)$ in the steady case. By applying the Laplace-Stieltjes transform and the methods of matrix theory, the steady distribution of an arbitrary demand is found to be

$$\bar{F}^*(x) = e^{B(x-\tau)} \bar{F}^*(\tau) + \int_{\tau}^x e^{B(x-y)} \bar{\varphi}(y) dy \quad x > \tau$$

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Unilinear system with ...

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$$B = \begin{pmatrix} \lambda & 0 & 0 & \dots & -\lambda \\ 0 & -\lambda & \lambda & \dots & 0 \\ \dots & \dots & \dots & \dots & \dots \\ 0 & \dots & \dots & -\lambda & \lambda \end{pmatrix}, \bar{\varphi}(x) = \begin{pmatrix} \varphi(x) \\ 0 \\ \vdots \\ 0 \end{pmatrix}, \bar{F}^* = \begin{pmatrix} F_1^*(x) \\ F_2^*(x) \\ \vdots \\ F_k^*(x) \end{pmatrix}$$

where

$$\varphi(x, \Pi(x)) = \int_0^{\tau} [1 - H(x - y)] dF_k^*(y)$$

$$\Pi(z) = \sum_{i=1}^k z^i F_i^*(0)$$

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Unilinear system with ...

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D251/D308

$P\{y_1 \leq x\} = H(x)$ is the distribution function of the service times
and λ is the expectation occurring in the formula for $G_k(x)$.

ASSOCIATION: Instytut matematyki AN URSS (Institute of Mathematics
of the AS UkrSSR) ✓

PRESENTED: by B. V. Hnyedenko, Academician

SUBMITTED: March 2, 1962

Card 4/4

BRODI, S.M. (Kiyev)

Limit theorem in the queueing theory. Ukr. mat. zhur. 15 no.1:
76-79 '63. (MIRA 16:3)

(Queueing theory)
(Limit theorems (Probability theory))

BRODICH, N. V.

Handbook for railroad make-up men and yard couplers. Izd. 2., ispr. i dop. Moskva, Gos. transp. zhel-dor. izd-vo, 1954. (Mic 55-3587) Collation of the original, as determined from the film: 354 p.

Microfilm Slavic 387 AC

1. Welding. 2. Railroads - Cars - Maintenance and repair

CZECHOSLOVAKIA / UNITED STATES

HYNIE, S.; KRISHNA, G.; ~~BRODIE, B.B.~~; Pharmacological Institute,
Medical Faculty, Charles University (Farmakologicky Ustav Fak.
Vseob. Lek. KU), Prague; Lab. Chem. Pharmacol. National Heart
Institute, NIH, Bethesda, Md.

"Mechanism of the Antilipolytic Effect of Betalytic DCI and
Alphalytic Phentolamine."

Prague, Ceskoslovenska Fysiologie, Vol 15, No 5, Sep 66, pp
410 - 411

Abstract: The degree of lipolysis is a function of the content of
the cyclic 3',5'-adenosine monophosphate in the fatty tissue.
The antilipolytic effect of DCI and of Phentolamine was tested
in vitro on rat fatty tissue. DCI in low concentrations blocks
only noradrenalin lipolysis; phentolamine in low concentrations
did not have an antilipolytic effect. At high concentrations
($10^{-3}M$) both drugs inhibit the activity of lipase. 3 Western ref-
erences. Submitted at 14 Days of Pharmacology at Smolenice, 16
Feb 66.

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CZECHOSLOVAKIA

DLABAC, A.; COSTA, E.; ~~BRODIE, B.B.~~; Research Institute of Pharmacy and Biochemistry (Vyzkumny Ustav pro Farmacii a Biochemii), Prague.

"Relationship Between the Central Effects of Reserpine and the Rate of Release of 5-Hydroxytryptamine."

Prague, Ceskoslovenska Fysiologie, Vol 15, No 5, Sep 66, p 418

Abstract: The correlation between the effect of reserpine and the decrease in the amount of 5-hydroxytryptamine (5-HT) in the homogenate of rat brain was investigated. Intravenous application of reserpine causes a fast exponential rate of decrease in the 5-HT content of the brain. The intensity of the central effect is proportional to the rate of initial reduction of 5-HT. A higher dose increases the intensity and the duration of the effects. Maximum rate of reduction in 5-HT was found after an administration of 5 and 10 mg/kg of reserpine. No references. Submitted at 14 Days of Pharmacology at Smolenice, 17 Feb 66.

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period
BRODIN, B. V., Cand Geol-Mineral Sci -- (diss) "Comparative
Characteristics of the Material Composition of Ore Deposits ~~in~~
the Kurgan Flexus ^U ~~of~~ Talas Abata^U." Mos, 1958. 16 pp. (Min
Higher Ed USSR, Mos Inst of Non-Ferrous Metals and Gold im M. I.
Kalinin^U, Chair of Mineralogy and Petrography), 110 copies.
(KL, 7-58, 109)

BRODIN, B.V.

Endogenetic goethite in the Ken-Shanyk ore deposit. Zap. Vses.
min. ob-va 87 no. 4:496-498 '58. (MIRA 12:1)
(Talas Ala-Tau--Goethite)

BRODIN, B.V.

Mineralogy of tin in ores of the Kurgan deposit. Zap. Kir. otd.
Vses. min. ob-va no.1:57-65 '59. (MIRA 14:3)
(Talas Ala-Tau—Tin)

BRODIN, B.V.

Mineral composition of ores in the Kurgan deposit (Talas Ala-Tau),
their formation and distribution. Geol. rud. mestorozh. no.5:57-73
S-O '59. (MIRA 13:2)

1. Institut tsvetnykh metallov im. M.I. Kalinina, Moskva.
(Talas Ala-Tau--Ore deposits)

BRODIN, B.V.

Tin minerals and their paragenesis in ore deposits of the
Kurgan group in the Talas Ala-Tau. Zap.Vses.min.ob-va 88
no.2:144-151 '59. (MIRA 12:8)
(Talas Ala-Tau--Tin)

BRODIN, B.V.

Myrmekite intergrowth of galena with chalcocite. Zap. Vses.
min. ob-va 89 no.4:415-423 '60. (MIRA 13:11)
(Myrmekites) (Galena) (Chalcocite)

BRODIN, B.V.

Wurtzite from the Ken-Shanyk deposit. Trudy Min. muz. no.11:
157-170 '61. (MIRA 16:7)

(Talas Ala-Tau—Wurtzite)

DYMKOV, Yu.M.; BRODIN, B.V.

Reddening of minerals in uranium-bearing veins. Atom. energ. 10
no.1:35-42 Ja '61. (MIRA 13:12)
(Uraninite) (Hematite)

BRODIN, B.V.; DYMKOV, Yu.M.

Montroseite from hydrothermal veins of the Pribram deposit. Zap.-
Vses.min.ob-va 90 no.6:653-659 '61. (MIRA 15:2)
(Pribram region--Montroseite)

BRODIN, B.V.

Internal structure and interrelations of nodular segregations
of some sulfides. Zap.Vses.min.ob-va 91 no.5:589-595 '62.

(MIRA 15:11)

(Sulfides)

BRODIN, B.V.

Role of diffusion in the replacement of minerals and the
decomposition of solid solutions. Trudy Min. muz. no.14:
79-95 '63. (MIRA 16:10)

(Mineralogical chemistry) (Solutions, Solid)

BRODIN, B.V.; DYMKOV, Yu.M.

Hard bitumens in uranium-bearing veins. Atom.energ. 16
no. 5:432-437 My '64. (MIRA 17:5)

~~BRODIN, I.S.~~

Automatic unit for testing gas meters with a rated discharge
up to 1.000 cubic meter per hour. Izm.tekh. no.5:53-57 My
'62.

(MIRA 15:6)

(Gas meters--Testing)

BRCEIN, I.S.

Gas rotational meters of the Ivano-Frankovsk Instrument Plant.
Gaz. prom. 9 no.2:19-20 '64. (MIRA 17:12)

L 9629-66

EWT(1)/EWA(h)/ETC(m)

WW

ACC NR: AP6000038

SOURCE CODE: UR/0115/65/000/010/0061/0062

AUTHOR: Brallov, E. S.; Brodin, I. S.; Sitnitskiy, Yu. I.; Chuchman, T. S.

ORG: None

TITLE: Improving the accuracy of a gas rotation meter

SOURCE: Izmeritel'naya tekhnika, no. 10, 1965, 61-62

TOPIC TAGS: gas flow, flow meter, measuring instrument, error

ABSTRACT: One of the main disadvantages of gas rotation meters, especially in research applications, is the considerable error ($\pm 2\%$) of industrially manufactured devices. The authors present the results of a study which is directed toward the improvement of the accuracy of such meters. The main cause of error is the overflow of gas through gaps between the moving rotors and the walls of the housing, depending on the pressure drop at the meter. The latter, in turn, depends on the flow rate. Hence, a mode of operation in which the pressure drop at the meter equals zero should eliminate or substantially reduce the error. In order to achieve this the rotors should be powered not by the energy obtained from the gas being measured, but from an external source. With this purpose, the authors designed and tested an automatic system which maintains zero pressure drop in a gas rotation meter (Fig. 1). Tests show that, in spite of large inertia of the rotors of a (RS-400) gas meter, the transfer process in the system does not exceed 30 sec. The work was per-

Card 1/2

UDC 681.122

L 9629-66

ACC NR: AP6000038

formed by L'vov Polytechnic Institute (L'vovskiy politekhnicheskiy institut) on orders from Ivano-Frankovskiy Instrument Building Plant (Ivano-Franskovskiy priborostroitel'nyy zavod). Orig. art. has: 2 figures.

(FD - ferrodynamic sensor; M - master; EA - electronic amplifier; RM - reversing motor; S - selsyn; MA - magnetic amplifier; AL - armature loop of d-c motor; DYA - dynamoelectric amplifier)

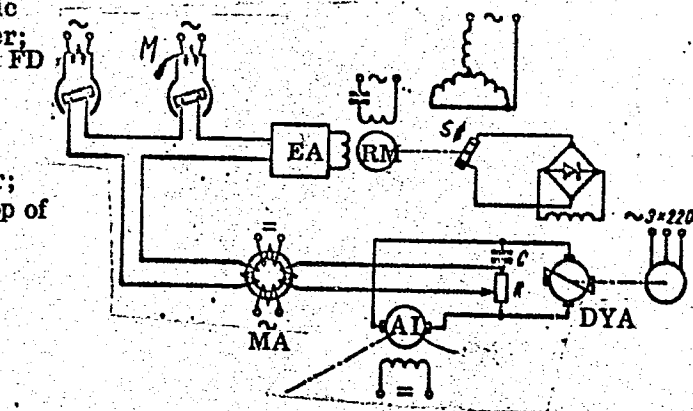


Fig. 1 - Automatic system for maintaining zero pressure drop.

SUB CODE: 14 / SUBM DATE: none

Card

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BRADIN, M. S.

Absorption and dispersion of light in stilbene crystals at room temperature and below. M. S. Bradin. *Optika i Spektroskopiya* 1, 878-80(1956). The absorption characteristics of cryst. plates (0.2-0.5 μ thick) of stilbene formed by sublimation onto quartz plates with optically parallel walls were investigated with an interferometer by a photographic method with polarized light. The absorption curves showed a shift in the absorption bands by ~ 100 cm⁻¹ in the direction of the long-wave region as the temp. of the crystal was gradually lowered from room temp. to 80° K. For the weak components the force of the oscillator (f) calcd. from the equation (given below) was found to be 0.13: $f = \frac{4\pi^2 N e^2}{m} \int \chi(\omega) d\omega$ (where ω = cyclic frequency corresponding to the max. in the absorption; m and e = electron mass and charge; N = no. of mols. per unit vol.). It remained practically const. throughout the entire range of temps. The dispersion curves in the region of transparency had a sharp increase in slope at the temp. of liquid N. The max. on the absorption curve and on the dispersion curve corresponded, resp., to each other. On the basis of the distribution of intensities in the two spectroscopic components and of the orientation of mols. in the lattice, the direction of the dipole moments transitions were approximated:

transition	polarization	γ (cm ⁻¹)	f
I	N_p	29600	0.13
	N_s	29900	0.64
II	N_p	40000	1.1
	N_s	40700	0.3

(Here, the N_s , N_p = polarization along the axis of the greatest and the smallest n , resp.; f = force of the oscillator for the k transition.) The first transition was polarized in the direction of the middle axis, the second in a direction close to the axis of CH=CH of the mol. — A. B. Kozlov

Inst. Physics, AS USSR

BRODIN, M.S.

Dichroism of some polycyclic crystals. Ukr. fiz. zhur. 1 no.4:
376-381 Q-D '56. (MLRA 10:2)

1. Institut fiziki AN URSS.
(Stilbene--Optical properties)
(Dichroism)

BRODIN, M.S.; MEDVEDEV, V.S.; PRIKHOT'KO, A.F.

Cryostats used in Jamin interferometry at liquid hydrogen temperatures. Prib.1 tekhn.eksp.no.3:96-98 M-D '56. (MLRA 10:2)

1. Institut fiziki AN USSR.
(Interferometry) (Cryostat)

dispersion curves the oscillator strengths of the 1st and 2nd
electron transition of the I crystal are 1.01, and also the ones
for the first transition of II. According to these results the
crystals of I and II absorb strongly and are dichroic. Thus
for the 1st electron transition in I the strength of the
linear polarization along the c axis is 0.99, and for II
it is 0.94. At the same time for the 2nd transition in II
the factor is 0.4.

W. J. L. J. J.

MT

BRODIN, M. S. Cand Phys-Math Sci -- (diss) "Study of the absorption and dispersion of light in crystals of certain polycyclic ~~compounds~~ compounds."
Kiev, 1957. 7 pp 20 cm. (Acad Sci UkrSSR. Inst of Physics), 100 copies. (KL, 13-57, 97)

PRIKHOT'KO, A.F.

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p.3

PHASE I BOOK EXPLOITATION 509/1365

L'vov. Universytet

Materialy X Vsesoyuznogo soveshchaniya po spektroskopii. t. 1: Molekulyarnaya spektroskopiya (Papers of the 10th All-Union Conference on Spectroscopy. Vol. 1: Molecular Spectroscopy) [L'vov] Izd-vo L'vovskogo univ-ta, 1957. 499 p. 4,000 copies printed. (Series: Its: Fizichnyy zbirnyk, vvp. 3/8/)

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Card 1/30

Brodin, M.S., and A.F. Prikhod'ko. Absorption and Dispersion of Light in Certain Molecular Crystals

16

Prikhod'ko, A.F., and M.T. Shpak. Polarization of Absorption Bands of Impurities in Crystals

21

Card 3/30

Inst. Physics, AN URSR

AUTHORS: Brodin, M.S. and Prikhot'ko, A.F.

51-4-7/25

TITLE: Dispersion and absorption of light in anthracene crystals at 20.4°K. (Dispersiya i pogloshcheniye sveta v kristallakh antratsena pri 20.4°K).

PERIODICAL: "Optika i Spektroskopiya" (Optics and Spectroscopy) 1957, Vol.2, No.4, pp.448-453 (U.S.S.R.)

ABSTRACT: Colourless anthracene crystals strongly absorb in the near ultraviolet at 24500-29000 cm^{-1} (first electron-vibration transition in the crystal) and also at 37000-42500 cm^{-1} (second transition). This paper reports on the refractive index dispersion and the absorption coefficients of anthracene crystals at 20.4°K at 25000-37000 cm^{-1} for two directions of the light vector: parallel and perpendicular to the monoclinic crystal axis. To study the dispersion a small Jamin interferometer was crossed by a spectrograph. The method is due to I.V.Obreimov (Jubilee collection of papers on the 70th birthday of Acad. A.F.Ioffe, p.523, 1950 - in Russian). Spectrograms were obtained with and without a crystal in the light beam. The absorption was measured with the cooperation of S.Z.Shul'ga (see also Zavodskaya laboratoriya, current number in print). A photomultiplier FEY-18 was used. For the dispersion and absorption measurements a quartz spectrograph of medium dispersion

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Dispersion and absorption of light in anthracene crystals
at 20.4°K. (Cont.) 51-4-7/25

was used. Low-pressure hydrogen lamps were used as light sources. Glan-Thompson prisms were used as polarizers. The iron arc spectrum was used as wavelength standard. The accuracy of measurement of the refractive index was about 2% and of the absorption coefficient 8-10%. The results are given in four graphs. The refractive index at 20.4°K oscillates strongly (from 1 to 6) at 25000-28000 cm⁻¹; its value outside this region is 1.5-2. The absorption coefficients (values up to 0.60) are shown for both 293°K and 20.4°K; oscillations here occur at 25 000 - 29 000 cm⁻¹. From the absorption coefficients and the refractive index the reflectivity curves (values up to 0.50) are constructed; again oscillations occur at 25 000 - 29 000 cm⁻¹. In all graphs the largest maxima occur at about 25 000 cm⁻¹. The oscillator strengths of electron-vibration transitions were calculated to be for the three directions: (a) parallel to the monoclinic axis, (b) perpendicular to that axis and (c) at right angles

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Dispersion and absorption of light in anthracene crystals^{51-4-7/25}
at 20.4°K. (Cont.)

to (a) and (b). Their values were found to be: (a) 0.12,
(b) 0.05 and (c) 0.13. There are six figures (including one
half-tone plate), and nine references, seven of which are
Slavic.

ASSOCIATION: Institute of Physics, Academy of Sciences of the
Ukrainian S.S.R., Kiev. (Institut Fiziki AN USSR, Kiev).
SUBMITTED: August 18, 1956.
AVAILABLE: Library of Congress

Card 3/3

BRODIN, M.S.; PRIKHOT'KO, A.F.

Absorption and scattering of light by various molecular crystals.
Fiz. sbor. no. 3:16-21 '57. (MIRA 11:8)

1. Institut fiziki AN USSR.
(Anthracene—Spectra)

BRODIN, M.S.; PRIKHOT'KO, A.F. [Prykhot'ko, A.F.]

Absorption and dispersion of light in certain crystals of the polycyclic series [In Ukrainian with summary in English]. Ukr. fiz. zhur. 3 no.1:79-87 Ja-F '58.

(MIRA 11:4)

1. Institut fiziki URSR.

(Stilbene--Optical properties)
(Acetylene--Optical)

AUTHORS: Brodin, M.S., Pakhomova, O.S. and Prikhod'ko, A.F. SOV/51-5-2-4/26
 TITLE: Absorption of Light by Stilbene Crystals at 20°K (Pogloshcheniye sveta kristallami stil'bena pri 20°K)
 PERIODICAL: Optika i Spektroskopiya, 1958, Vol 5, Nr 2, pp 123-127 (USSR)

ABSTRACT: The authors obtained and analysed the absorption spectrum of crystalline stilbene at 20°K. The crystals were prepared by sublimation and were attached to quartz glass plates. The spectra were photographed in polarized light using an ISP-22 quartz spectrograph. A hydrogen lamp with a "uviolet" window or a krypton lamp GSVD-120 were used as light sources. Iron spectrum was used for calibration. A FEU-18 photomultiplier was used as the receiver. Photographs of the two polarized components of the absorption spectrum (at 20°K) of a stilbene monocrystal, 0.2 μ thick, are shown in Fig 1. Fig 2 gives the absorption curve of a stilbene crystal at 20.4°K for vibrations parallel to the N_p axis. Fig 3 gives the absorption spectra of a stilbene crystal (curve 1) and a solid solution of stilbene in toluene (curve 2) at 20°K (vibrations parallel to the N_g axis). The authors make the following deductions from Fig 3. (1) The intensity of the first (purely electronic) band, as compared with the other bands, is considerably greater.

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Absorption of Light by Stilbene Crystals at 20°K

SOV/51-5-2-4/26

crystal spectrum than the intensity of the corresponding band in the solid solution spectrum. (2) 770 cm^{-1} vibration frequency is present in the pure crystal spectrum. The same vibration has a frequency of 745 cm^{-1} in the solid solution of stilbene in dibenzyl but it is absent in the solid solution of stilbene in toluene. (3) The bands corresponding to the harmonics of the 1590 cm^{-1} vibrations are stronger than the fundamental bands in the N_g -component of the pure crystal spectrum, but they are weaker in the solid solution spectra. The authors used crystals in optical contact with quartz plates or layers produced by melting between two quartz plates. In both cases cooling to the liquid-hydrogen temperature produced considerable mechanical strain due to the difference between the thermal expansion coefficients of quartz and stilbene. Such strains affect polarization of separate bands and their spectral position. The vibrational structure, however, is practically unaffected but the purely electronic bands are altered considerably and this has to be allowed for in making of any deductions. There are 3 figures and 5 Soviet references.

Card 2/2

ASSOCIATION: Institut fiziki, AN UkrSSR, g. Kiyev (Institute of Physics, Academy of Sciences of the Ukrainian S.S.R., Kiyev) 1. Stilbene crystals--Preparation
SUBMITTED: September 14, 1957 2. Single crystals--Spectrographic analysis

BRODIN, M.S.; PAKHOMOVA, O.S.; PRIKHOT'KO, A.F.

Absorption of light by stilbene crystals at 20°K. Opt. 1 spektr. 5
no. 2:123-127 Ag '58. (MIRA 11:10)

1. Institut fiziki AN YSSR, g. Kiyev.
(Stilbene)
(Absorption of light)
(Low temperature research)

24(2), 24(7)

SOV/48-22-11-8/33

AUTHORS:

Brodin, M. S., Soskin, M. S.

TITLE:

Connection of Absorption and Dispersion in Crystals at the Example of the 1,2-Benzanthrazen (Svyaz' pogloshcheniya i dispersii v kristallakh na primere 1,2-benzantratsena)

PERIODICAL:

Izvestiya Akademii nauk SSSR, Seriya fizicheskaya, 1958, Vol 22, Nr 11, pp 1316-1319 (USSR)

ABSTRACT:

In this work the absorption and dispersion spectra of a few molecular crystals have been examined together and the results have been compared to those of the Kramers' formula type. As object for an examination the 1,2-benzanthrazen-crystal was chosen. It was chosen over other crystals because it has two main-axes of the indicatrix of the indices of refraction in the plane of the lamellae produced in sublimation. For the comparison of the curves of dispersion and absorption the general integrals were used. They were found using Kramers' phenomenologic method. They can also be found on basis of the micro-theory (Ref 8). The experimental data which were used for the comparison are represented in figures 1 and 2. By comparing the sizes of the turning of these curves, which were determined experi-

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HOV/48-22-11-8/33

Connection of Absorption and Dispersion in Crystals at the Example of the
1,2-Benzanthrazen

mentally and calculated, an obvious divergence is shown. It is far beyond the limit of error (the calculated turning of the scale is at least three times smaller than the one determined experimentally). This has been observed on all 1,2-benzanthrazen-crystals examined. This divergence will be smaller for the same spectral-component at a normal temperature, but it will be still of importance. Similar phenomena have been observed earlier with crystals of other substances at low temperatures. There are 2 figures and 8 references, 6 of which are Soviet.

ASSOCIATION: Institut fiziki Akademii nauk USSR (Institute of Physics,
AS UkrSSR)

Card 2/2

SOV/51-6-1-5/30

AUTHORS: Brodin, M.S., Prikhod'ko, A.F. and Soskin, M.S.

TITLE: On Non-Validity of the Kramers--Kronig Dispersion Relationships in the Case of Molecular Crystals at Various Temperatures (O neachlydenii dispersionnykh sootnosheniy Kramersa--Kroniga v sluchaye molekulyarnykh kristallov pri razlichnykh temperaturakh)

PERIODICAL: Optika i Spektroskopiya, 1969, Vol 6, Nr 1, pp 28-32 (USSR)

ABSTRACT: Kramers--Kronig formulae (Ref 1) give the relationship between refractive indices and the absorption coefficients of a sample. They are derived from Maxwell's electromagnetic equations and should hold wherever Maxwell's theory holds. For some substances the Kramers--Kronig formulae were found to hold at room temperature (Refs 2, 3). The same substances when tested at low temperatures showed a disagreement between theory and experiment. The present paper describes an attempt at verification of these relationships in strongly absorbing molecular crystals of anthracene, 1,2-benzanthracene, stilbene and tolane which were studied at temperatures of 290, 80 and 20°K. The experimental data on absorption and dispersion of light in monocrystals of anthracene, stilbene and tolane were published earlier (Refs 5-7). The present paper gives the results obtained on 1,2-benzanthracene; these results

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On Non-Validity of the Kramers--Kronig Dispersion Relationships in the Case of
Molecular Crystals at Various Temperatures

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are shown in Fig 1 (absorption spectrum at 20°K) and Fig 2 (dispersion curve at 20°K - curve 1 was obtained experimentally and curve 2 was calculated). To measure the absorption and dispersion, thin samples, cut from monocrystals, were used. The monocrystals themselves were produced by sublimation. Measurements were made in polarized light. The dispersion curves were obtained interferometrically, by means of a Jamin interferometer. The absorption curves were measured photometrically using photographic film and photoelectric multipliers. The results reproduced from Refs 5-7 and those reported on 1,2-benzanthracene show that in some molecular crystals the Kramers--Kronig dispersion relationships may not be obeyed. The departure from these relationships is greatest at the lowest temperatures and it

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SOV/51-6-1-5/30

On Non-Validity of the Kramers--Kronig Dispersion Relationships in the Case of
Molecular Crystals at Various Temperatures

decreases with increase of temperature (about three times on increase of temperature from 20 to 290°K). The Kramers--Kronig formulae apply fully at room temperature in stilbene and tolane. On the other hand measurements on anthracene in light polarized parallel to the b-axis show that the Kramers--Kronig relationships are not obeyed even at room temperature. There are 2 figures and 10 references, 7 of which are Soviet, 2 English and 1 German.

SUBMITTED: March 29, 1958

Card 3/3

24(4), 24(6)

AUTHORS: Brodin, M.S. and Soskin, M.S.

SOV/51-6-5-7/34

TITLE: Studies of the Absorption Spectrum of 1,2-Benzanthracene Monocrystals in the Region of the Lowest Electron Transitions (Issledovaniye spektra pogloshcheniya monokristalla 1,2-benzantratsena v oblasti nizhayshikh elektronnykh perekhodov)

PERIODICAL: Optika i Spektroskopiya, 1959, Vol 6, Nr 5, pp 600-604 (USSR)

ABSTRACT: The authors obtained absorption spectra of 1,2-benzanthracene monocrystals (grown by sublimation) in polarized light at 20 and 290°K, in order to fill the gap in the spectral data on simple aromatic hydrocarbons (no work on the spectra of 1,2-benzanthracene was published so far). The absorption spectra were obtained for two components (a and b) in the region 25 000-33 000 cm⁻¹, which corresponds to the lowest singlet electron transitions. These spectra are shown in Figs 2 and 3, for 20 and 290°K respectively. 1 and 2 in Figs 2 and 3 denote the b-component and the a-component absorption curves respectively. The authors determined the integral intensities (areas S under the curves) of individual absorption bands and of the whole spectrum. The results of these calculations are given in Tables 1 (for 20°K) and 2 (for 290°K). The polarization ratio S_b/S_a (S_a , S_b are the corresponding areas under

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Studies of the Absorption Spectrum of 1,2-Benzanthracene Monocrystals in the Region
of the Lowest Electron Transitions

SOV/51-5-5-7/34

the a-component and b-component curves) and the "oriented gas model" were used to deduce the direction of the electric moment (polarization) of electron transitions in the 1,2-benzanthracene molecule. One of these transitions (1L_b) is directed along the y-axis of the molecule and the other (1L_a) makes an angle of 60° with that axis (Fig 1, diagram 2). The $27\,160\text{ cm}^{-1}$ frequency corresponds to the origin of the second transition. The authors investigated also the optical activity of the 1,2-benzanthracene crystals. Since the monocrystals were not sufficiently thick the results were only qualitative but optical activity was detected and the rotation constant in the region 3460 \AA was found to be $15\text{--}20\text{ deg/mm}$. Acknowledgment is made to A.F. Prihot'ko for his advice. There are 4 figures, 2 tables and 10 references, 4 of which are Soviet, 5 English and 1 mixed (Soviet and English).

SUBMITTED: May 12, 1963

Card 2/2

24(3), 24(4), 24(2)

SOV/51-7-1-12/27

AUTHORS: Brodin, M.S. and Lubchenko, A.F.

TITLE: The Effect of Temperature on Propagation of Electromagnetic Waves in Crystals in which Excitons are Formed (Vliyaniye temperatury na rasprostraneniye elektromagnitnykh voln v kristallakh, v kotorykh voznikayut eksitony)

PERIODICAL: Optika i spektroskopiya, 1959, Vol 7, Nr 1, pp 83-88 (USSR)

ABSTRACT: The authors discuss equations which give the refractive (n) and absorption (χ) indices for crystals in which excitons may be formed. These equations are given for weak coupling of excitons with the lattice vibrations. It is shown that with increase of temperature these equations tend to assume the form of the usual equations for n and χ in classical crystal optics. This tendency increases with the increase of the effective exciton mass and with the increase of the strength of the exciton-phonon interaction. Using Kramers--Kronig formulae the authors calculated the dispersion curves for anthracene and stilbene and compared the calculated curves with experimental ones (Figs 1, 2). It was found, in agreement with the above conclusions, that the differences between the calculated and experimental curves become

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The Effect of Temperature on Propagation of Electromagnetic Waves in Crystals in
which Excitons are Formed

SOV/51-7-1-12/27

smaller as temperature increases. At a given temperature the agreement between experimental and theoretical curves improves on transition from the electronic absorption bands to near electron-vibrational bands. Acknowledgments are made to A.F. Prihot'ko and A.S. Davydov for their advice. There are 2 figures, 1 table and 13 references, 11 of which are Soviet, 1 translation from German into Russian and 1 English.

SUBMITTED: July 23, 1958.

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24(7), 24(2)

AUTHORS: Brodin, M.S. and Prikhod'ko, A.F.

SO7/51-7-1-26/27

TITLE: The Effect of Thickness of Anthracene Crystals on their Absorption Curves at 20°K (Vliyanie tolshchiny kristallov antratsena na ikh krivyye pogloshcheniya pri temperature 20°K)

PERIODICAL: Optika i spektroskopiya, 1959, Vol 7, Nr 1, pp 132-133 (USSR)

ABSTRACT: Plane-parallel monocrystalline layers of anthracene, prepared by sublimation, were used. They were deposited on quartz plates and their optical density was measured photoelectrically, using a photomultiplier. The absorption coefficients were calculated by means of the usual formula [Eq (1)] in which $\lg(I_0/I)$ is the measured optical density corrected for losses by reflection and d is the crystal thickness. Since the thicknesses of the crystals were comparable with wavelengths used, a formula which is more exact than Eq (1) should be employed which would allow for interference in thin films. The authors found, however, that due to strong absorption by anthracene the error due to the use of Eq (1) was small and could be neglected. At 20°K the authors measured $\lg(I_0/I)$ as a function of frequency of about 20 crystals with thicknesses between 0.15 and 0.40 μ . The absorption coefficient curves of these crystals were found to depend on their

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The Effect of Thickness of Anthracene Crystals on their Absorption Curves at 20°K

thicknesses. By way of illustration the authors show in a figure on p 133 the absorption curves of the ϵ -component in the region 25000-27000 cm^{-1} for four crystals, 0.16, 0.21, 0.29 and 0.4 μ thick (curves 1, 2, 3, 4 respectively). The absorption curves obtained for the a -component of the spectrum showed a similar dependence on thickness. These results indicate a serious departure from the Lambert--Burger law and it is difficult to see the reason for this behaviour. The authors point out that this dependence on thickness was observed only at 20°K: the absorption curves measured at room temperature were found to be independent of thickness. The authors suggest that the reason for the low-temperature anomalies may lie in the fact that the exciton absorption formulae may not be valid at low temperatures. There are 1 figure and 7 references, 4 of which are Soviet and 3 English.

SUBMITTED: February 21, 1959

Card 2/2

AUTHORS: Brodin, M.S., Prikhod'ko, A.F. and Soskin, M.S.

SOV/51-7-2-25/34

TITLE: On Certain Dispersion Anomalies of Strongly Absorbing Crystals (O
nekotorykh osobennostyakh dispersii sil'no pogloshchayushchikh
kristallov)

PERIODICAL: Optika i spektroskopiya, 1959, Vol 7, Nr 2, pp 266-267 (USSR)

ABSTRACT: The authors showed earlier (Ref 1, 2) that certain molecular crystals, such as 1,2-benzanthracene and anthracene, do not obey the Kramers--Kronig dispersion relationships and that the degree of the departure from these relationships increases with lowering of temperature. The authors used a Jamin interferometer to obtain the dispersion curves throughout the region of the first electron-vibrational transition, with the exception of the wavelengths where maximum absorption occurred in strong bands, i.e. wavelengths at which anomalous dispersion took place. The present paper describes studies of these anomalous dispersion regions in thin crystals of 1,2-benzanthracene and anthracene. An arrow in Fig 1 shows the maximum of the first long-wavelength absorption band in the ϵ -component of the spectrum of 1,2-benzanthracene (0.25 μ thick) at 20°K. It is seen that the interference bands in the region of the arrow are apparently split into two and overlap, which is not predicted by the

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On Certain Dispersion Anomalies of Strongly Absorbing Crystals SOV/51-7-2-25/34

usual theory of anomalous dispersion. It is suggested that the observed effect is due to propagation of two identically polarized waves with unequal path differences. This dispersion peculiarity was not observed in 1,2-benzanthracene at room temperature (Fig 2). In the case of anthracene the splitting and overlapping of interference bands at the absorption maxima was observed both at low and at room temperatures. The two waves, which are not due to the usual birefringence, were recently predicted (Refs 5, 6) for the region of exciton absorption in crystals. The authors suggest that this may in fact explain the observed anomalies. If this interpretation is correct such anomalies should not be observed in impurity crystals. The latter conclusion is confirmed by a photograph of the interference system in the absorption region of anthracene molecules present in a crystal of dihydroanthracene (Fig 3); no anomalies are seen in that photograph. The authors point out that the observed anomalies occur only in the case when the Kramers--Kronig relationships are not obeyed. There are 3 figures and 6 references, 5 of which are Soviet and 1 English.

SUBMITTED: February 21, 1959

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S/181/60/002/009/017/036
B004/B056

26.1512
AUTHOR:

Brodin, M. S.

TITLE:

The Optical Properties of CdS Single Crystals

PERIODICAL: Fizika tverdogo tela, 1960, Vol. 2, No. 9, pp. 2152 - 2157

TEXT: It was the aim of the present paper to extend the measurement of the refractive index μ and the reflection coefficient R of CdS single crystals to the range of short wavelengths at 290 and 20°K, to investigate the anisotropy of the dispersion curves, and to check the Kramers - Kronig correlation. Single crystals having a thickness of from 5 to 1.5 μ and grown in the gaseous phase were used. The λ -dependence of the refractive indices was recorded by an interferometer according to a method described previously (Ref. 10). The intensity of the reflected light was measured by means of an apparatus described in Fig. 1 and compared with a standard made from quartz or aluminum mirror. A photomultiplier of the type ФЭУ-18 (FEU-18) was used as a receiver of the light energy. The values of the refractive indices were fairly stable in the case of various crystals within the region of transmissivity to visible light, ✓

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The Optical Properties of CdS Single Crystals S/181/60/002/009/017/036
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and the curves adapted themselves well to one another. Figs. 2,3 show the λ -dependence of the dispersion of a 2 μ thick crystal at the temperatures mentioned, which was measured in polarized light, and two polarization directions $\perp C$ and $\parallel C$, which are perpendicular to each other. The curve corresponding to the light vector $\perp C$ breaks off sooner in the violet region than the curve for $\parallel C$. The two curves intersect before reaching the absorption band. For 290°K the point of intersection is at $\lambda = 5080 \text{ \AA}$, for 20°K at 4880 \AA . At this point, the crystal is not birefringent, and behind it, birefringence changes its sign. Checking of the Kramers - Kronig correlation succeeded for the $\parallel C$ spectral component in the case of which the intensive absorption band was exceeded and the total dispersion amplitude within the band could be determined. Within the spectral band investigated (4750 - 5000 \AA) the experimental data were found to be in agreement with the Kramers - Kronig formula. In this way, the assumption of an exciton character of the absorption beginning at 5000 \AA is refuted. Recording of the reflection curve $R(\lambda)$ (Fig. 4) showed differences by far exceeding the experimental errors in various samples, which is attributed to the unstable properties of the crystal surface due to oxidation or excess of one of the two components of the

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The Optical Properties of CdS Single Crystals S/181/60/002/009/017/036
B004/B056

compound. The dichroism of the reflection repeats the course of the dispersion curve. The curves for $\parallel C$ and $\perp C$ intersect. The position of the point of intersection, however, changes in the individual samples. The author thanks A. F. Prikhod'ko for discussions. There are 4 figures and 14 references: 6 Soviet, 6 US, 1 Japanese, and 1 German.

ASSOCIATION: Institut fiziki AN USSR, Kiyev (Institute of Physics of the AS UkrSSR, Kiyev)

SUBMITTED: July 8, 1959

Card 3/3

BRODIN, M.S.; PRIKHOT'KO, A.F. [Prykhot'ko, A.F.]; SOSKIN, M.S.

Optical properties of crystals. Part 1. Ukr. fiz. zhur. 5 no.6:725-
743 N-D '60. (MIRA 14:3)

1. Institut fiziki AN USSR.
(Crystals--Optical properties)

BRODIN, M.S.; PEKAR, S.I.

Experimental proof of the existence of anomalous additional light waves in crystals in the exciton absorption region. Zhur. eksp. i teor. fiz. 38 no.1:74-81 Jan '60. (MIRA 14:9)

1. Institut fiziki Akademii nauk Ukrainskoy SSR.
(Anthracene crystals) (Excitons) (Light)

85701

S/056/60/038/006/044/049/XX
B006/B070

24.3500 (1035, 1114, 1138)

AUTHORS: Brodin, M. S., Pekar, S. I.

TITLE: Additional Anomalous Light Waves in Anthracene in the Region
of Exciton Absorption

PERIODICAL: Zhurnal eksperimental'noy i teoreticheskoy fiziki, 1960,
Vol. 38, No. 6, pp. 1910 - 1912

TEXT: The existence of additional anomalous light waves in crystals in the region of exciton absorption was theoretically predicted by Pekar (Refs. 1-3). An experimental proof of this assumption was suggested by the present authors in Ref.4. In the same paper also data of previous measurements of light absorption in anthracene plates of different thickness had been evaluated (in the region of characteristic absorption, maximum at $25,200 \text{ cm}^{-1}$). It was found that the intensity of transmitted light as a function of the plate thickness was subject to fluctuations. In the present "Letter to the Editor", a report is made on the absolute measurement of the intensity of light by the method of photographic

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Additional Anomalous Light Waves in
Anthracene in the Region of Exciton
Absorption

S/056/60/038/006/044/049/XX
B006/B070

photometry (reference line: $24,720 \text{ cm}^{-1}$). A spectrograph of the type $\Delta\phi C-3$ (DFS-3) was used for 3 different thicknesses of the crystal. The measured optical density of the crystal as a function of its thickness for the frequency $25,108 \text{ cm}^{-1}$ is shown in a diagram ($T = 20^\circ\text{K}$). The oscillation character is here clearer than in the curves of Ref.4. It is noted that the separations Δl of the extrema are about equal (abscissas of the maxima: $l = 0.072, 0.128, 0.188, 0.245 \mu$; separations: $\Delta l = 0.056, 0.060, 0.057 \mu$; minima: $l = 0.105, 0.168, 0.233 \mu$; separations: $\Delta l = 0.063$ and 0.055μ). With this, the average period of oscillation is found to be 0.058μ , and the corresponding difference in the refractive indices of two interfering waves to be 6.9. The character of the curve shown in the Fig. is that of a theoretical curve describing the interference between two parallel anomalous waves polarized in parallel. These more accurate measurements again demonstrate the existence of additional anomalous waves in the crystal of anthracene, as is required by the theory. A. F. Prihot'ko is thanked for interest and discussions, and S. V. Marisova for help in experiments. I. V. Obreimov is mentioned.

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Additional Anomalous Light Waves in Anthracene in the Region of Exciton Absorption

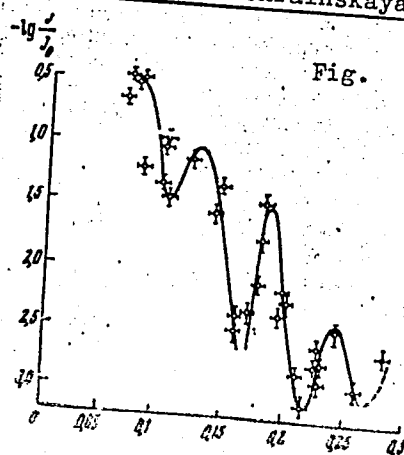
S/056/60/038/006/044/049/XX
B006/B070

There are 1 figure and 6 Soviet references.

ASSOCIATION: Institut Fiziki Akademii nauk Ukrainskoy SSR (Institute of Physics of the Academy of Sciences Ukrainskaya SSR)

SUBMITTED: March 19, 1959

Fig.



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S/185/61/006/005/016/019
D274/D303

AUTHORS: Brodin, M.S., and Krochuk, A.S.

TITLE: Fine structure and temperature dependence of CuCl
single crystals absorption-spectrum

PERIODICAL: Ukrayins'kyy fizychnyy zhurnal, v. 6, no. 5, 1961,
706 - 709

TEXT: Plane-parallel single-crystals, 5 - 40 μ thick, were prepared; no thinner plate could be obtained by mechanical means. The absorption spectra were obtained at temperatures of 4.2, 20.4, 150 and 290°K on a spectrograph with a dispersion of 4 Å/mm; the reflection spectra were photographed at 20.4°K with a dispersion of 10 Å/mm. A figure shows the curves of the absorption edge for a specimen 23 μ thick, at the four indicated values of the temperature. Two strong absorption-bands were observed, one at $T = 4.2^{\circ}\text{K}$, $\nu \approx 25834 \text{ cm}^{-1}$, and the other at $T = 20.4^{\circ}\text{K}$ with a reflection maximum at $\nu = 26435 \text{ cm}^{-1}$. These bands are apparently exciton bands. They were also observed in polycrystalline films by R. Reiss and

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Fine structure and temperature ...

S/185/61/006/005/016/019
D274/D303

S. Nikitine (Ref. 3: C.r. Acad. Sci., 250, 2862, 1960), where the presence of weak bands between the above two bands, was observed. The authors were however unable to observe these weak bands. On the other hand, the authors observed two weak and very narrow bands on the longwave side of the band $\nu = 25834 \text{ cm}^{-1}$. They are best observed at liquid helium temperature, and are broadened on increase of temperature. These bands have two interesting properties: their relative intensity varies from crystal to crystal, so that in one specimen the first band is stronger and in another specimen - the second; in addition, their intensity depends little on thickness of specimen. Another characteristic feature of the absorption bands of CuCl , is the non-monotonic temperature shift of the absorption edge; the authors concluded that the maximum of the strong absorption band $\nu = 25834 \text{ cm}^{-1}$ at $T = 4.2^\circ\text{K}$, is also non-monotonically temperature dependent. The narrow bands, 25642 and 25694 cm^{-1} have a similar temperature shift. The non-monotonic character of the temperature shift is an indication of the complex absorption mechanism and deserves further study. Further, the peculiar features of the narrow bands are related to the inhomogeneity of impurity centers,

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whose field influence the corresponding exciton levels. As the authors did not conduct crystallographic investigations, they were unable to ascertain the relationship between the anisotropic character of the crystals and the peculiarities of the narrow bands. There are 2 figures and 6 references: 3 Soviet-bloc and 3 non-Soviet-bloc. The reference to the English-language publication reads as follows: R. Coelho, Techn. Rep. 143, Mass (USA), October 1960.

ASSOCIATION: Instytut fizyki AN URSR m. Kyiv (Institute of Physics AS UkrSSR, Kyiv)

SUBMITTED: June 10, 1961

✓

Card 3/3

BRODIN, M.S.; MARISOVA, S.V.

Davydov splitting and polarization ratio in the region of the
first electron transition of an anthracene crystal. Opt. i
spektr. 10 no.4:473-476 Ap '61. (MIRA 14:3)
(Anthracene crystals—Spectra)

26.2532

28104
S/181/61/003/009/038/039
B108/B138

AUTHORS: Brodin, M. S., Vitrikhovskiy, N. I., Strashnikova, M. I.

TITLE: Structure of the spectra of $\text{CdS}_x\text{CdSe}_{1-x}$ and $\text{CdS}_x\text{ZnS}_{1-x}$ hybrid crystals at 20°K

PERIODICAL: Fizika tverdogo tela, v. 3, no. 9, 1961, 2882-2885

TEXT: Ye. F. Gross and V. V. Sobolev (DAN SSSR, 133, 56, 1960) have shown that the emission spectrum of CdS-CdSe solid solutions at low temperatures bears the same character as the CdS spectrum. V. V. Yeremenko (FTT, II, 2602, 1960) studied the low-temperature absorption spectra of $\text{CdS}_x\text{CdSe}_{1-x}$ hybrids but could not find any fine structure, apparently because he did not have sufficiently thin specimens at his disposal. In order to elucidate this problem, and to obtain data on the character of the excitation in pure crystals the authors studied 10 - 20 μ thick hexagonal CdS-CdSe and CdS-ZnS hybrids. The back-reflection spectra were taken by means of a Hilger-E2 spectrograph. Photomicrographs of such spectra

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Structure of the spectra of CdS_x ... S/181/61/003/009/038/039
B108/B138

taken at a temperature of 20°K from CdS , $\text{CdS}_{0.95}\text{CdSe}_{0.05}$, and $\text{CdS}_{0.94}\text{ZnS}_{0.06}$ single crystals are shown in the Figure (a, b, and c, respectively). It was found that the absorption spectrum of any CdSe.CdS hybrid has a fine structure. The bands 1 and 2 in the figure are shifted to the longwave side by about 60 cm^{-1} when the CdSe concentration in CdS rises by one percent by weight, whereas the band 3 is shifted only by about 30 cm^{-1} . The back reflection in polarized light indicates that the absorption of $\text{CdS}_x\text{CdSe}_{1-x}$ has a distinct dichroism: The $\perp\text{C}$ absorption edge is shifted to longer waves. The $\text{CdS}_x\text{CdSe}_{1-x}$ hybrids exhibit an inversion point of the refractive index at which the crystal foils are not birefringent. The back-reflection and, consequently, the absorption spectra of $\text{CdS}_x\text{ZnS}_{1-x}$ hybrids are very similar to those of pure CdS , particularly when the ZnS concentration is low. When the ZnS concentration is higher than about 20%, the back reflection bands become more and more blurred. This phenomenon is explained by the interference bands arising when the rays are reflected in the transparent region from both the front and the back surface of the

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crystal foil. The argument that the crystal is inhomogeneous is not justified since then interference bands could never exist. The difference in the band shift with concentration is explained as follows: The optical electrons of CdS responsible for the bands 1 and 2 in the valence band are preferably connected with the Cd ion, the electron causing the band no. 3, however, with the S ion. There are 1 figure and 8 references: 5 Soviet and 3 non-Soviet. The two references to English-language publications read as follows: S. Y. Czyzak et al., J. Opt. Soc., 47, 240, 1957. D. G. Thomas, Y. Y. Hopfield. Phys. Rev., 116, 573, 1959.

ASSOCIATION: Institute fiziki AN USSR (Physics Institute of the AS UkrSSR) Institut poluprovodnikov AN USSR Kiyev (Semiconductor Institute of the AS UkrSSR Kiyev)

SUBMITTED: May 22, 1961

Card 3/4

BRODIN, M.S.; KROCHUK, A.S.

Fine structure and temperature dependence of the edge of the
absorption spectrum of CuCl monocrystals. Ukr. fiz. zhur.
6 no.5:706-709 S-0 '61. (MIRA 14:11)

1. Institut fiziki AN USSR, g. Kiyev.
(Copperchloride crystals--Spectra)

BABENKO, V.P.; BRODIN, M.S.; SOSKIN, M.S.

Cryostat for dispersion measurements at low temperatures.
Prib. i tekhn. eksp. 6 no. 6: 140-141 N-D '61. (MIRA 14:11)

1. Institut fiziki AN USSR.
(Cryostat)

BRODIN, M.S.; MARISOVA, S.V.

Spectral distribution of the dependence of the transparency of anthracene single crystals on their thickness. Ukr.fiz.shur. 6 no.6:745-750 N-D '61. (MIRA 16:5)

1. Institut fiziki AN UkrSSR, Kiyev.
(Anthracene crystals—Optical properties)

S/051/61/011/006/005/012
E039/E385

AUTHORS: Brodin, M.S. and Dovgiy, Ya.O.

TITLE: Optical properties of mixed crystals. I.
Characteristic absorption in dispersive mixtures of
anthracene in crystals of 9,10-dihydroanthracene

PERIODICAL: Optika i spektroskopiya, v.11, no.6, 1961, 742-749

TEXT: The authors studied the optical properties of mixed crystals and the dependence of these properties on the concentration of the mixture. Absorption curves were obtained for dispersive mixtures of anthracene in crystals of 9,10-dihydroanthracene. The samples were grown by a sublimation method, producing monocrystalline plane parallel plates of various thickness, suitable for quantitative optical measurement. The absorption measurements were obtained with a photographic recording large dispersion spectrograph (4 Å/mm). When calculating the absorption coefficients corrections were made for the spectral intensity distribution of the source, the spectral sensitivity of the plate and, for important measurements, the reflection of light from the crystal faces. The polarization measurements were
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Optical properties of

S/051/61/011/006/005/012
E039/E385

made using Nicol prisms and the dispersion curves were measured by an interferometric method. The absorption curves were measured in the region of the first electron transition of anthracene for a series of samples containing an admixture of various concentrations in the range 0.2 - 5%. Measurements were carried out at several temperatures (290, 145, 77 and 20 °K) but the principal investigation was at 20 °K. A typical absorption curve for 0.6% anthracene in 9,10-dihydroanthracene at 20 °K, showing also the effect of polarized light, is given in Fig. 1a, where \circ - light polarized parallel to b axis, Δ - light polarized parallel to a axis, κ - absorption coefficient and ν - frequency. The experimental data on dispersion were compared with the Kramer-Kronig relationship. There is a small disagreement at low concentrations which, according to the measurements, grows with increasing concentration of the mixture. If it is assumed that the anthracene molecules are uniformly distributed, the distances between them would be approximately 20 Å but the investigation described here shows that the spacings between the anthracene molecules are unequal. Acknowledgments are expressed

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S/051/61/011/006/005/012
EO59/E385

to A.F. Prikhod'ko for his interest in the present work.
There are 2 figures, 2 tables and 8 references: 6 Soviet-bloc
and 2 non-Soviet-bloc. The English-language references
mentioned are: Ref. 3: D.S. McClure, J. Chem. Phys., 22, 1668,
1954; 24, 1, 1956; Y.W. Sidman, Phys. Rev., 102, 96, 1956;
Y.W. Sidman, J. Chem. Phys., 25, 115, 1956; Y.W. Sidman,
D.S. McClure - J. Chem. Phys., 24, 757, 1956;
Ref. 5: Y. Iball - J. Chem. Soc., 1074, 1938.

SUBMITTED: January 2, 1961

Card 3/4
3

24.7000

40884

S/181/62/004/009/018/045
B108/B186

AUTHORS: Brodin, M. S., and Strashnikova, M. I.

TITLE: Singularities in the dispersion and shape of the exciton absorption bands of CdS single crystals

PERIODICAL: Fizika tverdogo tela, v. 4, no. 9, 1962, 2454 - 2460

TEXT: A homogeneous layer (0.095 μ) of CdS was studied. The absorption curves were measured at 77, 20, and 4°K with a grating spectrograph. The dispersion curves were determined by a method described earlier (PTE, no. 3, 96, 1956; no. 6, 140, 1961). Two principal absorption bands were found (Fig. 1) with linearly polarized light ($E \perp C$ and $E \parallel C$; C is the hexagonal crystal axis). With $E \perp C$, the band A has a greater dispersion than the band B although its peak height and integral intensity are nearly the same as those of B (Fig. 3). This makes the Kramers-Kronig dispersion relation no longer applicable in this section of the spectrum. It is concluded that the exciton whose excitation leads to the band B enters into no strong interaction with the phonons. There are 4 figures and 2 tables.

ASSOCIATION: Institut fiziki AN USSR, Kiev (Physics Institute AS UkrSSR, ~~Soviet Union~~ Kiev)

Submitted April 1962

S/185/62/007/001/004/01:
D299/D302

AUTHORS: Brodin, M.S., and Dovhyy, Ya.O.

TITLE: On some peculiar features of the exciton-absorption spectrum of sodium-uranylacetate single-crystals

PERIODICAL: Ukrayins'kyy fizychnyy zhurnal, v. 7, no. 1, 1962, 30 - 36

TEXT: The spectra were analyzed in circularly-polarized light. The structure of the absorption bands and their intensity were determined. The dependence of the absorption on specimen thickness was studied. The investigations were carried out at low temperatures (20 and 4°K respectively). Absorption spectra of both dextro- and levorotatory single-crystals were obtained. At 20°K, nine narrow absorption bands were observed. At 4°K, each of these absorption bands separates into 2 - 3 components. The 9 absorption bands are characterized by a high degree of circular dichroism. These bands were not observed in the spectrum of a chilled solution (at 77°K) of sodium-uranylacetate. This fact in conjunction with the high degree of circular dichroism is characteristic of the dextro- and levorotatory single-crystals of sodium-uranylacetate. ✓
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On some peculiar features of the ...

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cular dichroism of the absorption bands, shows that the observed bands are of exciton type. A figure shows the absorption curves

$K(\nu) = \lg \frac{I_0}{I} / 4\pi d \nu \cdot 0.43$, at 20°K , for a dextrorotatory crystal, 0.525 mm thick. These curves are a measure of the intensity distribution in some of more characteristic narrow bands, as well as of the degree of circular dichroism. The absorption curves for the levorotatory crystals are almost identical with those for the dextrorotatory ones. Further, the dependence of the intensity of the absorption band (with frequency $\nu_m = 21135 \text{ cm}^{-1}$), on specimen thickness, was in-

vestigated; the thickness varied between 170 - 650 μ . For 3 special points (21133, 21134.2, 21137 cm^{-1}), a non-monotonic dependence of the optical density on thickness was observed. The shape and the position of the maximum of the band vary with specimen thickness. These facts might be related to the presence of anomalous light waves in the absorption band under consideration or to quadrupole origin of the band; it is also possible that both factors are responsible. A final explanation would require further optical and crystallogra-

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On some peculiar features of the ...

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D299/D302

phic investigations. There are 5 figures and 8 references: 7 Soviet-bloc and 1 non-Soviet-bloc.

ASSOCIATION: Instytut fizyky Am URSR (Institute of Physics of the AS UkrRSR), Kyiv

SUBMITTED: March 6, 1961

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38849

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S/185/62/007/006/014/014
D407/D301

AUTHORS: Brodin, M. S. and Strashnikova, M. I.

TITLE: Absorption and dispersion of light in CdS single-crystals at 20°K

PERIODICAL: Ukrayins'kyy fizychnyy zhurnal, v. 7, no. 6, 1962, 681-683

TEXT: The absorption and dispersion curves of thin CdS single-crystals in the special region 20 500 - 21 500 cm^{-1} were measured. This region contains two intensive absorption bands. The present investigation constitutes a continuation based on more adequate data, of an earlier study by M. S. Brodin (Ref. 9: FTT, 2, 2152, 1960), which had the purpose of verifying the Kramers-Kronig dispersion relations. CdS single-crystals (0.1 μ thick) were fixed on quartz plates. The measurements were conducted at 20°K in polarized light. A spectrograph with a dispersion of 4 $\text{\AA}/\text{mm}$ was used. The light intensity was measured by the method of photographic photom-etering. The dispersion curves were obtained by the interference

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Absorption and dispersion ...

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method. The position of the maximum of one of the bands is noted as a peculiar feature of the spectrum. The two intensive bands are denoted by A and B. By comparing the spectra a and b (corresponding to two different angles of light polarization) it was found that on turning the crystal the maximum of the band A is shifted towards the long-wavelength side; a similar shift was observed if the crystal rotated about its vertical axis. No corresponding shift of the band B was observed. The shift of the band A cannot be explained by variations in reflection, neither by crystal inhomogeneities nor by its rotation. A satisfactory interpretation of the observed shift requires further experimental work. It is noted, however, that energy changes of the exciton transition as a function of the direction of polarization and of the wave vector were predicted by theory. From a figure it is evident that the band A ($\nu_m = 20\,567\text{ cm}^{-1}$) is completely polarized in a direction normal to the C-axis. At $T = 20^\circ\text{K}$, it has no fine structure. The maximum of the band B is slightly shifted as a result of the polarization. The bands A and B have practically equal intensity. The dispersion

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Absorption and dispersion ...

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D407/D301

curves, just as the absorption curves, were obtained at two different polarization angles and also at normal incidence. A maximum of the dispersion curve was observed in the region of the first band, followed by an anomalous section and a sharp drop. The authors were unable to sufficiently reduce the amplitude of the dispersion curves. In the region of the band B it was possible to construct the entire dispersion curve. The character of the dispersion curves for the ordinary and extraordinary waves is an indication of the complexity of the double refraction of the crystal in the absorption region; the crystal has inversion points of the refraction index in the entire spectral range. On comparing the absorption and dispersion curves, it is noted that although the bands A and B are similar in many respects, yet they are related to quite dissimilar amplitudes of the dispersion curves. The Kramers-Kronig relations are not completely corroborated by the above experimental results. There are 3 figures. f

ASSOCIATION: Instytut fizyki AN UkrRSR, Kyiv (Institute of Physics of the AS UkrRSR, Kiev)

SUBMITTED: March 2, 1962

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S/051/62/012/002/017/020
E202/E192

AUTHORS: Brodin, M.S., and Dovgiy, Ya.O.

TITLE: Experimental study of circular dichroism and optical activity of sodium uranyl acetate single crystals

PERIODICAL: Optika i spektroskopiya, v.12, no.2, 1962, 285-290

TEXT: This paper opens an extensive study of the above single crystals, which due to their optical activity are likely to display the effects of space dispersion. The absorption curves of crystal origin bands were measured in detail to determine their polarisation and intensity. The luminescence spectrum was also taken and briefly commented on. Rectangular prism shaped, highly polished crystals of 0.2-1 mm thickness were grown from pure solution. l- and d-rotatory specimens were studied. Circle polarised light was used and the sample was kept in a cryostat within 77-20 °K. Absorption curves were measured photographically using high dispersion spectrograph (2 and 4 Å/mm), and a three-lens illuminator for the slit. Spectra were photographed through a stepwise intensity wedge, analysed with a microphotometer and the values used to determine the coefficient of absorption.
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Experimental study of circular ...

S/051/62/012/002/017/020
E202/E192

To determine the amount and spectral distribution of the optical activity, transmission spectra of crystals placed between crossed polarisers were taken. It was found that the absorption spectrum of a single crystal, as well as a frozen solution consisted of strong, periodically reappearing groups of bands, which were attributed to the excitation of molecular electrons. In addition, the single crystal spectrum had seven characteristically spaced narrow bands with considerable circular dichroism. These were present chiefly only in a spectrum of one direction of circular polarisation, while the opposite direction of polarisation led to very weak intensities. The distribution of optical activity showed that the crystal rotates the plane of polarisation only in the vicinity of bands with a pronounced circular dichroism. There are 3 figures.

SUBMITTED: January 21, 1961

Card 2/2

BRODIN, M.S.

Study of the form of absorption bands in tolan single crystals
Ukr.fiz.zhur. 7 no.4:417-421 Ap '62. (MIRA 15:8)

1. Institut fiziki AN UkrSSR, g. Kiyev.
(Acetylene) (Absorption spectra)

45077

S/051/63/014/001/014/031
E039/E120

24.3950

AUTHORS: Brodin, M.S., and Krochuk, A.S.

TITLE: Peculiarities of the optical absorption of CuCl single crystals

PERIODICAL: Optika i spektroskopiya, v.14, no.1, 1963, 88-93

TEXT: Samples of CuCl, which belongs to the cubic system and has T_d^2 symmetry, were prepared in the form of thin parallel plates (10 μ or more thickness) from large single crystals grown at the Institut Kristallografii AN SSSR (Institute of Crystallography, AS USSR). These crystals were uniform and colourless with a mirror finish on their surfaces. They were mounted either in paper holders or on quartz plates; the method of fixing did not influence their spectra. Photographic recording was used on a diffraction grating spectrograph with a dispersion of 4 $\text{\AA}/\text{mm}$. A Xenon lamp was used as a light source; this provided a continuous spectrum in the region investigated. Absorption curves were obtained at temperatures of 290, 150, 77, 20.4 and 4.2 $^{\circ}\text{K}$. Reflection spectra were obtained using normal incidence at 290, 150, 77 and 20.4 $^{\circ}\text{K}$. In the latter case wedge-

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Peculiarities of the optical ...

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shaped samples were used to avoid light reflected from the back surface entering the spectrograph. At low temperatures strong bands were observed which are ascribed to exciton excitation. The temperature displacement of the edge of the spectrum is not monotonic, hence confirming the complex character of absorption. Near the intensity maximum $25\,830\text{ cm}^{-1}$ at 4.2°K three narrow weak bands were observed at $25\,642$, $25\,694$ and $25\,706\text{ cm}^{-1}$ which can be grouped in a hydrogen-like series, described by the relation

$$\nu_n = 25\,710 - \frac{68}{n^2}\text{ cm}^{-1}$$

where $n = 1, 2$ and 3 .
There are 4 figures.

SUBMITTED: December 20, 1961

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